

Product Datasheet

Anti-Mad3 (Human) Antibody FL550 Conjugate



Overview

Catalog # 75-250-FL550

Conjugate FL550 Ex: 550 nm, Em: 575 nm

 $\begin{tabular}{lll} Isotype & IgG2b \\ Clone Number & N129/15 \\ Size & 200 \ \mu L \\ Concentration & 0.5 \ mg/mL \\ \end{tabular}$

Host Species Mouse Monoclonal

Format Purified by Protein A chromatography

Buffer PBS with 0.09% azide

Applications ICC
Species Reactivity Human

Immunogen Fusion protein amino acids 116-206 (C-terminus) of human Mad3 (accession number Q9BW11)

produced recombinantly in E. Coli

Molecular Weight 25 kDa

Cite this Antibody Antibodies Inc Cat# 75-250-FL550, RRID: AB 2939909

Details

Target Description MAX Dimerization Protein 3 is encoded by the gene MXD3. MXD3 is a member of the Myc

superfamily. MXD3 is a transcriptional repressor which forms a heterodimer with the cofactor MAX to form a sequence-specific DNA-binding protein complex which recognizes the core sequence 5'-CAC[GA]TG-3'. MXD3 is expressed in lung and blood. Diseases associated with MXD3 include

Perianal Hematoma and Bartholin's Duct Cyst.

Specificity Does not cross-react with Mxi1

Purification Method Produced by in vitro bioreactor culture of hybridoma line followed by Protein A affinity

chromatography and conjugation of purified mAb. Purified mAbs are >90% specific antibody.

Quality Control Tests Each new lot of antibody is quality control tested on cells overexpressing target protein and

confirmed to give the expected staining pattern.

Storage Aliquot and store at \leq -20°C for long term storage. For short term storage, store at 2-8°C. For

maximum recovery of product, centrifuge the vial prior to removing the cap.

Our Guarantee

As an original manufacturer, we are dedicated to creating quality and reproducible antibodies that further your research. We provide personalized customer support from the scientists that made the antibody and offer a free replacement or 100% refund if we cannot resolve an issue. Order today and experience our 50+ year passion for science.

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